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MEMORANDUM

TO:

Mr. Addison Rice

Anderson, Mulholland and Associates

DATE: January 8, 2015

FROM: R. Infante

FILE: 1412151A

RE:

Data Validation

Air samples SDG: 1412151A

SUMMARY

Full validation was performed on the data for several gas samples analyzed for selected volatile organic compounds by method Compendium Method TO-15: Determination Of Volatile Organic Compounds (VOCs) In Air Collected In Specially-Prepared Canisters And Analyzed By Gas Chromatography/Mass Spectrometry (GC/MS), January, 1999. The samples were collected at the Bristol Myer Squib-Building 5 VI facility, Humacao, PR site on December 08-09, 2014 and submitted to Eurofins Air Toxics, Inc. of Folson, California that analyzed and reported the results under delivery group (SDG) 1412151A.

The sample results were assessed according to USEPA data validation guidance documents in the following order of precedence: Compendium Method TO-15. Determination Of Volatile Organic Compounds (VOCs) In Air Collected In Specially-Prepared Canisters And Analyzed By Gas Chromatography/Mass Spectrometry (GC/MS), January, 1999; Validating Air Samples. Volatile Organic Analysis of Ambient Air in Canisters by Method TO-15, (SOP # HW-31. Revision #4. October, 2006 The QC criteria and data validation actions listed on the data review worksheets are from the primary guidance document, unless otherwise noted.

In general the data is valid as reported and may be used for decision making purposes. The data results are acceptable for use. Detected results for Toluene, 2-Propanol, and Acetone in samples 1412151-01A, 1412151-02A, and 1412151-03A exceed the instrument calibration range and are considered estimated values (E).

SAMPLES

The samples included in the review are listed below

Client Sample ID	Lab. Sample ID	Collected Date	Matrix	Analysis
BSIA-5 (2014)	1412151A-01A	12/10/2014	Air	VOCs
BSIA-3D (2014)	1412151A-02A	12/10/2014	Air	VOCs
BSIA-3 (2014)	1412151A-03A	12/10/2014	Air	VOCs
BSIA-11 (2014)	1412151A-04A	12/11/2014	Air	VOCs
BSIA-9 (2014)	1412151A-05A	12/11/2014	Air	VOCs

REVIEW ELEMENTS

Sample data were reviewed for the following parameters, where applicable to the method

- o Agreement of analysis conducted with chain of custody (COC) form
- o Holding time and sample preservation
- Gas chromatography/mass spectrometry (GC/MS) tunes
- o Initial and continuing calibrations
- o Method blanks/trip blanks/field blank
- o Canister cleaning certification criteria
- Surrogate spike recovery
- o Internal standard performance and retention times
- o Field duplicate results
- o Laboratory control sample/laboratory control sample duplicate (LCS/LCSD) results
- o Quantitation limits and sample results

DISCUSSION

Agreement of Analysis Conducted with COC Request

Sample reports corresponded to the analytical request designated on the chain-of-custody form.

Holding Times and Sample Preservation

Sample preservation was acceptable.

Samples analyzed within method recommended holding time.

GC/MS Tunes

The frequency and abundance of bromofluorobenzene (BFB) tunes were within the QC acceptance criteria. All samples were analyzed within the tuning criteria associated with the method.

Initial and Continuing Calibrations

VOCs (Method TO-15)

The percent relative standard deviations (%RSDs) and response factors (RFs) of all target analytes were within the QC acceptance criteria in the initial calibration. Correlation coefficients (r²) of target analytes were within the QC acceptance criteria. Ongoing accuracy of the instrument was determined by the analysis of a continuing calibration standard.

Method Blank/Trip Blank/Field Blank

Target analytes were not detected in laboratory method blanks for VOCs.

Summa canister met cleaning certification criteria.

Surrogate Spike Recovery

The surrogate recoveries were within the laboratory QC acceptance limits in all samples analyzed.

Internal Standard Performance

VOCs

Samples were spiked with the method specified internal standard. Internal standard are performance and retention times met the QC acceptance criteria in all sample analyses and calibration standards.

Laboratory/Field Duplicate Results

VOCs

Field/laboratory duplicates were analyzed as part of this data set. Target analytes meet the RPD performance criteria of +25 %.

LCS/LCSD Results

VOCs

LCS/LCSD (blank spike) were analyzed by the laboratory associated with this data package. Recoveries and RPD within laboratory control limits.

Quantitation Limits and Sample Results

Dilution was performed on samples BSIA-5 (2014) and BSIA-3 (2014) due to the presence of high level target species.

Detected results for Toluene, 2-Propanol, and Acetone in samples 1412151A-01A; 1412151A-02A; and 1412151A-03A exceed the instrument calibration range and are considered estimated values.

Calculations were spot checked.

Certification

The following samples 1412151A-01A; 1412151A-02A; 1412151A-03A; 1412151A-04A; and 1412151A-05A were analyzed following standard procedures accepted by regulatory agencies. The quality control requirements met the methods criteria except in the occasions described in this document. The results are valid. Some of the results were qualified.

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Rafael Infante

Chemist License 1888



Client Sample ID: BSIA-5 (2014) Lab ID#: 1412151A-01A

MODIFIED EPA METHOD TO-15 GC/MS FULL SCAN

File Name:		Date of Collection: 12/8/14-3:55:00 PM
Dil. Factor:	7.90	Date of Analysis: 12/16/14 03:27 PM

Compound	Rpt. Limit (ppbv)	Amount (ppbv)	Rpt. Limit (ug/m3)	Amount (ug/m3)
Benzene	0.79	Not Detected	2.5	Not Detected
Ethyl Benzene	0.79	47	3.4	200
m,p-Xylene	0.79	160	3.4	710
o-Xylene	0.79	15	3.4	63
Toluene	0.79	340 E	3.0	1300 E
2-Propanol	4.0	410 E	9.7	1000 E
4-Methyl-2-pentanone	0.79	160	3.2	660
Acetone	4.0	590 E	9.4	1400 E

E = Exceeds instrument calibration range.

Container Type: 6 Liter Summa Canister (100% Certified)

Surrogates	%Recovery	Method Limits
1,2-Dichloroethane-d4	101	70-130
Toluene-d8	105	70-130
4-Bromofluorobenzene	94 SOCIALO	70-130





Client Sample ID: BSIA-3D (2014) Lab ID#: 1412151A-02A

MODIFIED EPA METHOD TO-15 GC/MS FULL SCAN

File Name: Dil. Factor:	v121612 2.87	Date of Collection: 12/9/14 9:30:00 AM
DII. Factor.	2.01	Date of Analysis: 12/16/14 04:13 PM

Compound	Rpt. Limit (ppbv)	Amount (ppbv)	Rpt. Limit (ug/m3)	Amount (ug/m3)
Benzene	0.29	Not Detected	0.92	Not Detected
Ethyl Benzene	0.29	22	1.2	95
m,p-Xylene	0.29	78	1.2	340
o-Xylene	0.29	7.1	1.2	31
Toluene	0.29	150 E	1.1	570 E
2-Propanol	1.4	170 E	3.5	410 E
4-Methyl-2-pentanone	0.29	79	1.2	320
Acetone	1.4	270 E	3.4	650 E

E = Exceeds instrument calibration range.

Container Type: 6 Liter Summa Canister (100% Certified)

		Method	
Surrogates	%Recovery	Limits	
1,2-Dichloroethane-d4	105	70-130	
Toluene-d8	107	70-130	
4-Bromofluorobenzene	95	70-130	

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Client Sample ID: BSIA-3 (2014) Lab ID#: 1412151A-03A

MODIFIED EPA METHOD TO-15 GC/MS FULL SCAN

File Name: Dil. Factor:	v121613	Date of Collection: 12/9/14 9:20:00 AM
Dil. Factor.	2.98	Date of Analysis: 12/16/14 04:52 PM

Rpt. Limit (ppbv)	Amount (ppbv)	Rpt. Limit (ug/m3)	Amount (ug/m3)
0.30	Not Detected	0.95	Not Detected
0.30	28	1.3	120
0.30	98	1.3	420
0.30	9.2	1.3	40
0.30	190 E	1.1	710 E
1.5	200 E	3.7	500 E
0.30	100	1.2	410
1.5	320 E	3.5	750 E
	(ppbv) 0.30 0.30 0.30 0.30 0.30 1.5 0.30	(ppbv) (ppbv) 0.30 Not Detected 0.30 28 0.30 98 0.30 9.2 0.30 190 E 1.5 200 E 0.30 100	(ppbv) (ppbv) (ug/m3) 0.30 Not Detected 0.95 0.30 28 1.3 0.30 98 1.3 0.30 9.2 1.3 0.30 190 E 1.1 1.5 200 E 3.7 0.30 100 1.2

E = Exceeds instrument calibration range.

Container Type: 6 Liter Summa Canister (100% Certified)

Surrogates	%Recovery	Limits
1,2-Dichloroethane-d4	99	70-130
Toluene-d8	110	70-130
4-Bromofluorobenzene	102	70-130





Client Sample ID: BSIA-11 (2014) Lab ID#: 1412151A-04A

MODIFIED EPA METHOD TO-15 GC/MS FULL SCAN

File Name:	v121614	Date of Collection: 12/9/14 10:46:00 AM
Dil. Factor:	1.83	Date of Analysis: 12/16/14 05:37 PM

Compound	Rpt. Limit (ppbv)	Amount (ppbv)	Rpt. Limit (ug/m3)	Amount (ug/m3)
Benzene	0.18	Not Detected	0.58	Not Detected
Ethyl Benzene	0.18	2.5	0.79	11
m,p-Xylene	0.18	8.9	0.79	39
o-Xylene	0.18	1.0	0.79	4.3
Toluene	0.18	19	0.69	71
2-Propanol	0.92	28	2.2	70
4-Methyl-2-pentanone	0.18	9.0	0.75	37
Acetone	0.92	39	2.2	92

Container Type: 6 Liter Summa Canister (100% Certified)

		Method
Surrogates	%Recovery	Limits
1,2-Dichloroethane-d4	112	70-130
Toluene-d8	106	70-130
4-Bromofluorobenzene	97	70-130





Acetone

Air Toxics

0.80

Client Sample ID: BSIA-9 (2014) Lab ID#: 1412151A-05A

MODIFIED EPA METHOD TO-15 GC/MS FULL SCAN

File Name: Dil. Factor:	v121615 1.61		Date of Collection: 12/9/14 10:50:00 AM Date of Analysis: 12/16/14 06:14 PM			
Compound	Rpt. Limit (ppbv)	Amount (ppbv)	Rpt. Limit (ug/m3)	Amount (ug/m3)		
Benzene	0.16	Not Detected	0.51	Not Detected		
Ethyl Benzene	0.16	2.6	0.70	11		
m,p-Xylene	0.16	9.3	0.70	40		
o-Xylene	0.16	0.93	0.70	4.0		
Toluene	0.16	19	0.61	72		
2-Propanol	0.80	28	2.0	70		
4-Methyl-2-pentanone	0.16	8.8	0.66	36		

Container Type: 6 Liter Summa Canister (100% Certified)

		Method		
Surrogates	%Recovery	Limits		
1,2-Dichloroethane-d4	112	70-130		
Toluene-d8	104	70-130		
4-Bromofluorobenzene	94	70-130		

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Sample Transportation Notice
Relinquishing signature on this document indicates that sample is being shipped in compliance with all applicable local, State, Federal, national, and international laws, regulations and ordinances of any kind. Air Toxics Limited assumes no liability with respect to the collection, handling or shipping of these samples. Relinquishing signature also indicates agreement to hold harmless, defend, and indemnify Air Toxics Limited against any claim, demand, or action, of any kind, related to the collection, handling, or shipping of samples. D.O.T. Hotline (800) 467-4922

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Lab I.D. Field Sample I.D. (Location)	Can #	_	ate liection	Time of Collection	Analyses Reques	sted	Initial	final	Ssure/Vac	Stanosana en
OIA BSIA-5(2014)	34723	12/8	1/14	1555	see note :		<i>3</i> 0+	5		Final (per)
024 BSTA-30(2014)	35255			0930			30+	16		
084 BSTA - 3 (2017)	34450	12/0	1/14	0920	+1		30 ⁺	3.5		
044 BSTA-11 (2014)	1651	12/0	,/14	1046	11	•	30°	8.5		
054 BSIA-9 (ROL4)	34440	12/9	/19	1080			28	5		
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	Project Number:1412151A
	Date:12/08-09/2014
REVIEW OF VOLATILE ORGATINE following guidelines for evaluating volatile organics was actions. This document will assist the reviewer in using prodecision and in better serving the needs of the data users. The USEPA data validation guidance documents in the follows the company that the company	ere created to delineate required validation ofessional judgment to make more informed ne sample results were assessed according to ing order of precedence: QC criteria from
"Compendium Method TO-15. Determination of Volatile Org Specially-Prepared Canisters and Analyzed By Gas Chi January, 1999"; USEPA Hazardous Waste Support Branc Analysis of Ambient Air in Canisters by Method TO-15, (SOP QC criteria and data validation actions listed on the data revied document, unless otherwise noted. The hardcopied (laboratory name) _Eurofins	romatography/Mass Spectrometry (GC/MS), h. Validating Air Samples. Volatile Organic # HW-31. Revision #4. October, 2006). The ew worksheets are from the primary guidance
reviewed and the quality control and performance data summa	arized. The data review for VOCs included:
Lab. Project/SDG No.:1412151A No. of Samples:5	Sample matrix:Air
Trip blank No.: Field blank No.: Equipment blank No.: Field duplicate No.:1412151A-02A/1412151A-03A	
X Data CompletenessX Holding TimesX GC/MS TuningX Internal Standard PerformanceX BlanksX Surrogate RecoveriesN/A_ Matrix Spike/Matrix Spike Duplicate	X Laboratory Control SpikesX Field DuplicatesX CalibrationsX Compound IdentificationsX Compound QuantitationX Quantitation Limits
Overall Comments:_Selected_VOC's_by_method_TO-15	5
Definition of Qualifiers:	
J- Estimated results	
U- Compound not detected	
R- Rejected data UJ- Estimated nondetect	
OJ- ESUMALEO NONGELECT	
Reviewer: Calaul alaut	
Date:01/08/20 15	

DATA COMPLETENESS

MISSING INFORMATION	DATE LAB. CONTACTED	DATE RECEIVED
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All criteria were metX	
Criteria were not met	
and/or see below	

HOLDING TIMES

The objective of this parameter is to ascertain the validity of the results based on the holding time of the sample from time of collection to the time of analysis.

Complete table for all samples and note the analysis and/or preservation not within criteria

SAMPLE ID	DATE SAMPLED	DATE ANALYZED	pН	ACTION
	All samples analyzed w	ithin the recommended	method	holding time
	<u> </u>			
		·		

Criteria

Aqueous samples – 14 days from sample collection for preserved samples (pH \leq 2, 4°C), no air bubbles.

Aqueous samples – 7 days from sample collection for unpreserved samples, 4°C, no air bubbles. Soil samples- 7 days from sample collection.

Cooler temperature (Criteria: 4 + 2 °C): N/A - summa canisters

Actions

If the VOCs vial(s) have air bubbles, estimate positive results (J) and reject nondetects (R). If the % solids of soil samples is 10-50%, estimates positive results (J) and nondetects (UJ) If the % solid of soil samples is < 10%, estimate positive results (J) and reject nondetects (R). If holding times are exceeded but < 14 days beyond criteria, estimate positive results (J) and nondetects (UJ).

If holding times are exceeded but < 28 days beyond criteria, estimate positive results (J) and reject nondetects (R).

If holding times are grossly exceeded (> 28 days beyond criteria), reject all results (R).

If samples were not iced or if the ice were melted (> 10°C), estimate positive results (J) and nondetects (UJ).

		Criteri	All criteria were metX a were not met see below
GC/MS TUNING			
The assessment standard tuning (_	determine if the sample instrum	entation is within the
XThe BFB	performance results were	reviewed and found to be within th	ne specified criteria.
XBFB tuni	ng was performed for every	24 hours of sample analysis.	
If no, use profes qualified or reject		ine whether the associated data	should be accepted,
List	the	samples	affected:
If mass calibratio	n is in error, all associated	data are rejected.	

All criteria were metX_	
Criteria were not met	
and/or see below	

CALIBRATION VERIFICATION

Compliance requirements for satisfactory instrument calibration are established to ensure that the instrument is capable of producing and maintaining acceptable quantitative data.

Date of initial calibration:	10/16/14
Dates of continuing calibratio	n:12/16/14
Instrument ID numbers:I	MSD-V
Matrix/Level:	Air/low

DATE	LAB ID#	FILE	CRITERIA OUT RFs, %RSD, %D, r	COMPOUND	SAMPLES AFFECTED
			rations meet method s requirements.	pecific requirements. Initia	I calibration retention

Criteria

All RFs must be > 0.05 regardless of method requirements for SPCC.

All %RSD must be < 15 % regardless of method requirements for CCC.

All %Ds must be \leq 30% regardless of method requirements for CCC.

Method TO-15 does not specify criterion for the curve correlation coefficient (r). A limit for r of \geq 0.995 has therefore been utilized as professional judgment.

Actions

If any compound has an initial RF or a continuing RF of < 0.05, estimate positive results (J) and reject nondetects (R), regardless of method requirements.

If any compound has a %RSD > 15%, estimate positive results (J) and use professional judgment to qualify nondetects.

If any compound has a %RSD > 90%, estimate positive results (J) and reject nondetects (R).

If any compound has a % D > 30%, estimate positive results (J) and reject nondetects (R).

If any compound has a % D > 30%, estimate positive results (J) and nondetects (UJ).

If any compound has a % D > 90%, estimate positive results (J) and reject nondetects (R).

If any compound has r > 0.995, estimate positive results and nondetects.

A separate worksheet should be filled for each initial curve

All criteria were metX
Criteria were not met
and/or see below

V A. BLANK ANALYSIS RESULTS (Sections 1 & 2)

The assessment of the blank analysis results is to determine the existence and magnitude of contamination problems. The criteria for evaluation of blanks apply only to blanks associated with the samples, including trip, equipment, and laboratory blanks. If problems with any blanks exist, all data associated with the case must be carefully evaluated to determine whether or not there is an inherent variability in the data for the case, or if the problem is an isolated occurrence not affecting other data.

List the contamination in the blanks below. High and low levels blanks must be treated separately.

Laboratory blanks

DATE ANALYZED	LAB ID	LEVEL/ Matrix	COMPOUND	CONCENTRATION UNITS
All_metho	d_blank_meeth	_method_speci		
Summa_c			ation_criteria	
Field <u>/</u> Equipmen			· · · · · · · · · · · · · · · · · · ·	
DATE ANALYZED	LAB ID	LEVEL/ MATRIX	COMPOUND	CONCENTRATION UNITS
No_field/trip/eq	uipment_blanks	_analyzed_with	this_data_package	
		······································		
,		······································		
		14 - 110 Test, 1 - 1	The second secon	

All criteria were metX
Criteria were not met
and/or see below

VB. BLANK ANALYSIS RESULTS (Section 3)

Blank Actions

Action Levels (ALs) should be based upon the highest concentration of contaminant determined in any blank. Do not qualify any blank with another blank. The ALs for samples which have been diluted should be corrected for the sample dilution factor and/or % moisture, where applicable. No positive sample results should be reported unless the concentration of the compound in the samples exceeds the ALs:

ALs = 10x the amount of common contaminants (methylene chloride, acetone, 2-butanone, and toluene)

ALs = 5x for any other compounds

Specific actions are as follows:

If the concentration is < sample quantitation limit (SQL) and \le AL, report the compound as not detected (U) at the SQL.

If the concentration is \geq SQL but \leq AL, report the compound as not detected (U) at the reported concentration.

If the concentration is \geq SQL and > AL, report the concentration unqualified.

Notes:

High and low level blanks must be treated separately Compounds qualified "U" for blank contamination are still considered "hits" when qualifying for calibration criteria.

CONTAMINATION SOURCE/LEVEL	COMPOUND	CONC/UNITS	AL/UNITS	SQL	AFFECTED SAMPLES
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					· · · · · · · · · · · · · · · · · · ·
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17.6 4.					

All criteria were metX
Criteria were not met
and/or see below

SURROGATE SPIKE RECOVERIES

Laboratory performance of individual samples is established by evaluation of surrogate spike recoveries. All samples are spiked with surrogate compounds prior to sample analysis. The accuracy of the analysis is measured by the surrogate percent recovery. Since the effects of the sample matrix are frequently outside the control of the laboratory and may present relatively unique problems, the validation of data is frequently subjective and demands analytical experience and professional judgment.

List the percent recoveries (%Rs) which do not meet the criteria for surrogate recovery.

Matrix: solid/aqueous

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SURROGATE COMPOUND

ACTION

1,2-DICHLOROETHANE- Toluene- 4-BFE d4 d8

_Surrogate_recoveries_within_laboratory_control_limits					
QC Limits* (Air)					
LL_to_UL70to_130	_70to_13070to_130				

- * QC limits are laboratory in-house performance criteria, LL = lower limit, UL = upper limit.
- * If QC limits are not available, use limits of 80 120 % for aqueous and 70 130 % for solid samples.

Actions:

QUALITY	%R < 10%	%R = 10% - LL	%R > UL
Positive results	J	J	J
Nondetects results	R	ΠΊ	Accept

Surrogate action should be applied:

If one or more surrogate in the VOC fraction is out of specification, but has a recovery of > 10%.

If any one surrogate in a fraction shows < 10 % recovery.

All criteria were met
Criteria were not met
and/or see belowN/A

VII. A MATRIX SPIKE/MATRIX SPIKE DUPLICATE (MS/MSD)

This data is generated to determine long term precision and accuracy in the analytical method for various matrices. This data alone cannot be used to evaluate the precision and accuracy of individual samples. If any % R in the MS or MSD falls outside the designated range, the reviewer should determine if there are matrix effects, i.e. LCS data are within the QC limits but MS/MSD data are outside QC limit.

1. MS/MSD Recoveries and Precision Criteria

The laboratory should use one MS and a duplicate analysis of an unspiked field sample if target analytes are expected in the sample. If target analytes are not expected, MS/MSD should be analyzed.

AS OR MSD	COMPOUND	% R	RPD	QC LIMITS	ACTION
	_are_not_required_as			TO-15;_blank_sp	ike_used_to_assess

Actions:

QUALITY	%R < LL	%R > UL
Positive results	J	J
Nondetects results	R	Accept

MS/MSD criteria apply only to the unspiked sample, its dilutions, and the associated MS/MSD samples:

If the % R for the affected compounds were < LL (or 70 %), qualify positive results (J) and nondetects (UJ).

If the % R for the affected compounds were > UL (or 130 %), only qualify positive results (J).

If 25 % or more of all MS/MSD %R were < LL (or 70 %) or if two or more MS/MSD %Rs were < 10%, qualify all positive results (J) and reject nondetects (R).

A separate worksheet should be used for each MS/MSD pair.

All criteria were met
Criteria were not met
and/or see belowN/A

VII. B MATRIX SPIKE/MATRIX SPIKE DUPLICATE

MS/MSD - Unspiked Compounds

It should be noted that Method TO-15 does not specify a MS/MSD criteria for the unspiked compounds in the sample. A %RSD of < 50% has therefore been utilized as professional judgment.

If all target analytes were spiked in the MS/MSD, this review element is not applicable.

List the %RSD of the compounds which do not meet the criteria.

Sample ID:			Matrix/Level/Unit:			
COMPOUND	SAMPLE CONC.	MS CONC.	MSD CONC.	% RSD	ACTION	
				· · · · · · · · · · · · · · · · · · ·		
						
		······				
					······	
			-			

Actions:

^{*} If the % RSD > 50, qualify the positive result in the unspiked samples as estimated (J).

^{*} If the % RSD is not calculated (NC) due to nondetected value, use professional judgment to qualify the data.

All criteria were metX
Criteria were not met
and/or see below

VIII. LABORATORY CONTROL SAMPLE (LCS) ANALYSIS

This data is generated to determine accuracy of the analytical method for various matrices.

1. LCS Recoveries Criteria

LOOID

Where LCS spiked with the same analyte at the same concentrations as the MS/MSD? Yes or No. If no make note in data review memo.

List the %R of compounds which do not meet the criteria

	rc2 in	COMPOUND	% R	QC LIMIT
LCS/LCS	D_(Blank_spike	e)_analyzed_in_this_data_	package,_recoveries_ar	nd_RPD_within
laborator	y_control_limits.			
				· · · · · · · · · · · · · · · · · · ·
	· · · · · · · · · · · · · · · · · · ·			

- * QC limits are laboratory in-house performance criteria, LL = lower limit, UL = upper limit.
- * If QC limits are not available, use limits of 70 130 %.

Actions:

QUALITY	%R < LL	%R > UL
Positive results	J	J
Nondetects results	R	Accept

All analytes in the associated sample results are qualified for the following criteria.

If 25 % of the LCS recoveries were < LL (or 70 %), qualify all positive results (j) and reject nondetects (R).

If two or more LCS were below 10 %, qualify all positive results as (J) and reject nondetects (R).

2. Frequency Criteria:

Where LCS analyzed at the required frequency and for each matrix? <u>Yes</u> or No. If no, the data may be affected. Use professional judgment to determine the severity of the effect and qualify data accordingly. Discuss any actions below and list the samples affected.

			All criteria were met Criteria were not met and/or see belowN/A		
IX.	LABORATOR	Y DUPLICATE PRECISION			
	Sample IDs:	_1412151A-02A/1412151A-02A	Matrix:Air		

Field duplicates samples may be taken and analyzed as an indication of overall precision. These analyses measure both field and lab precision; therefore, the results may have more variability than laboratory duplicates which only laboratory performance. It is also expected that soil duplicate results will have a greater variance than water matrices due to difficulties associated with collecting identical field duplicate samples.

The project QAPP should be reviewed for project-specific information. Suggested criteria: RPD \pm 25% for air samples. If both samples and duplicate are <5 SQL, the RPD criteria is doubled.

SQL	SAMPLE CONC.	DUPLICATE CONC.	RPD	ACTION
RPD v	vithin the met	hod performand	e criter	ia.
		CONC.	CONC. CONC.	

Actions:

Qualify as estimated positive results (J) and nondetects (UJ) for the compound that exceeded the above criteria. For organics, only the sample and duplicate will be qualified.

If an RPD cannot be calculated because one or both of the sample results is not detected, the following actions apply:

If one sample result is not detected and the other is greater than 5x the SQL qualify (J/UJ).

If one sample value is not detected and the other is greater than 5x the SQL and the SQLs for the sample and duplicate are significantly different, use professional judgment to determine if qualification is appropriate.

If one sample value is not detected and the other is less than 5x, use professional judgment to determine if qualification is appropriate.

If both sample and duplicate results are not detected, no action is needed.

All criteria were metX
Criteria were not met
and/or see below

X. INTERNAL STANDARD PERFORMANCE

The assessment of the internal standard (IS) parameter is used to assist the data reviewer in determining the condition of the analytical instrumentation.

List the internal standard area of samples which do not meet the criteria.

- * Area of +40% or -40% of the IS area in the associated calibration standard.
- * Retention time (RT) within \pm 0.06 seconds of the IS area in the associated calibration standard.

DATE	SAMPLE ID	IS OUT	IS AREA	ACCEPTABLE RANGE	ACTION
	tandard_area_and_reation_standards		within_laboratory	_control_limits_for_	_both_samples
Actions:					

1. IS actions should be applied to the compound quantitated with the out-of-control ISs

QUALITY	IS AREA < -40%	IS AREA > + 40%
Positive results	J	J
Nondetected results	R	ACCEPT

If a IS retention time varies more than 0.330 seconds, the chromatographic profile for that sample must be examined to determine if any false positive or negative exists. For shifts of a large magnitude, the reviewer may consider partial or total rejection of the data for the sample fraction.

All criteria were metX	
Criteria were not met	
and/or see below	

XII. SAMPLE QUANTITATION

The sample quantitation evaluation is to verify laboratory quantitation results. In the space below, please show a minimum of one sample calculation:

1412151A-01A

Toluene

RF = 1.62872

[] = (6014361)(5.0)/(424896)(1.62872)

= 43.45 ppbv OK

All criteria were metX
Criteria were not met
and/or see below

XII.	OI	IANTI'	TATIOI	N I	INAL	ΓS
AH.	W.	<i>1</i> 771311	-		LILVII	

A. Dilution performed

	الله من		
ļ	Percent S	Colids	
'	COOM	ondo	
į	List samp	les which have ≤ 50	% solids
. •			
-			
-			